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C252 C28X C30Y C32Y C321 C322 C323 C327 C34Y
C341 C354 C366 C367 C385 C45Y C456 C51X C51Y
C510 C516 C535 C578 C61X C620 C626 C628 C630
C650 C657 C658 C670 C699 C70X C727 C740 C759
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None

(58) Field of Search

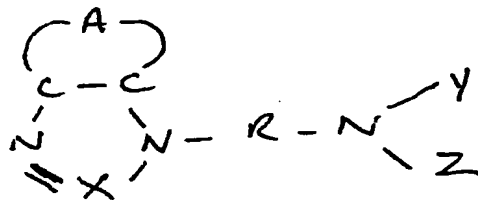
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(54) Blowing agent inhibitors for printing inks

(57) Printing ink comprises as a blowing agent inhibitor, benzimidazole or a compound of the formula



A= benzene, naphthene or cycloaliphatic ring,

X= N or C - R⁷ (R⁷ = H or alkyl)

R= C₁₋₅alkylene

Y= H, OH or organic moiety

Z= organic moiety

Y and Z together may form an organic ring.

A subset of these compounds is claimed per se (see claim 12).

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EMBOSSING INHIBITOR

The invention relates to blowing agent inhibitors and their use. In particular, the invention is directed to highly insoluble azoles, especially benzotriazole and benzimidazole

5 derivatives, which are effective blowing agent inhibitors.

The inhibitors are substantially insoluble in both water and alcohol and are capable of being ground and dispersed in situ in an ink composition.

In the present invention, azole derivatives such as
10 benzotriazole (BTA), tolyltriazole (TTA) and benzimidazole derivatives have been found to be crystalline solids of very high melting point and unusually low solubility in many solvents, including water and isopropyl alcohol. These derivatives can be readily micronized and dispersed into aqueous inks of widely varying
15 composition with no adverse impact on the stability of the ink or its printing and drying characteristics. Further, because of their very low solubility, these derivatives can be dispersed into typical ink compositions and ground in situ without adverse effects on the ink composition.

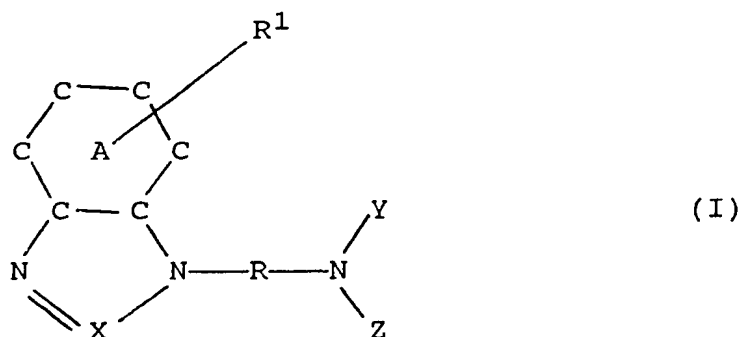
20 Relative to BTA, TTA and other aminomethyltriazole derivatives, the present inhibitors are also significantly less prone to undesirable post-printing migration, a problem commonly encountered in the process of chemical embossing. The inhibitors of the prior art diffuse, typically upward from
25 the printed surface into the bottom of the adjacent layer of foamable plastic substrate when a continuous sheet is tightly

wound and stored before further processing. This inhibitor contact-migration results in formation of faint images in the non-embossed areas of the surface, a phenomenon commonly referred to as "ghost embossing".

5 Accordingly, an object of the present invention is to provide an inhibitor for water-based inks which is universally compatible, does not destabilize the ink, dries without any tack, embosses satisfactorily and shows significantly reduced ghosting characteristics.

10 The term "azole" as used herein includes benzotriazole, tolyltriazole, naphthotriazole, cycloaliphatic triazole, benzimidazole, tolylimidazole, naphthimidazole, cycloaliphatic imidazole and their derivatives. Preferred compounds are those which have a room temperature aqueous solubility of less than
15 0.1% by weight or a room temperature isopropyl alcohol solubility/ of less than 5% by weight. These compounds do not interfere with the ink stability or drying characteristics of the ink composition. Therefore, the liquid ink has excellent shelf-life and dries without becoming tacky.

20 It is also an object of the present invention to provide a printing ink composition comprising a resin, solvent, and an inhibitor; the inhibitor being benzimidazole or a compound having the general formula



wherein the A ring is benzenoid, naphthenoid or saturated cycloaliphatic, the A ring being unsubstituted or substituted with R¹ which is an alkyl group of 1 to 4 carbon atoms, X being a nitrogen atom or the

group, wherein R⁷ is a hydrogen atom or an alkyl group of 1 to 4 carbon atoms, R is an alkylene group of 1 to 5 carbon atoms, Y is a hydrogen atom, a hydroxide group or an organic moiety and Z is an organic moiety or Y and Z when taken together with the nitrogen to which they are attached form an organic ring structure, the inhibitor when other than benzimidazole having a 24 hour room temperature isopropyl alcohol solubility of less than 5% by weight.

Preferably, the inhibitor has a 24 hour room temperature aqueous solubility of less than 0.1% by weight.

(II)

$$\begin{array}{c} =\text{C}-\text{R}^7 \\ | \\ \text{---} \end{array}$$

- 4 -

2,4,6-tris(benzotriazol-1'-yl methyl)-s-triazine,
2,4-bis(benzotriazol-1'-yl methyl) benzoguanamine,
N,N-bis(benzotriazol-1-yl methyl) glycine,
N-(benzotriazol-1-yl methyl)-4'-carboxybenzene
sulfonamide, N,N'-bis(benzotriazol-1-yl methyl)
naphthalene-1,5-disulfonamide (i.e. the compound prepared
in Example 24), N,N'-bis(benzotriazol-1-yl methyl)
benzene-1,3-disulfonamide (i.e. the compound prepared in
Example 25), 1-bis(benzotriazol-1'-yl methyl)-2-benzoyl
hydrazide, bis(benzotriazol-1-yl methyl) amine and
1,3-bis(5'-butyl benzotriazol-1'-yl methyl) urea. The
remaining undefined substituents on the saturated D ring
are hydrogen atoms. Preferably the A' and A'' rings are
the same and R¹ and R² are the same. R⁵ and R⁶ are
preferably the same, more preferably they are both
methylene groups. It is also preferred that X' and X''
are both nitrogen atoms.

A further object of the invention is to provide a
method of embossing a heat-foamable resinous material by
applying the printing ink composition of the present
invention to selected areas of the surface of a heat-
foamable resinous material, which material contains a
blowing agent, and subsequently heating the material to
or above the decomposition temperature of the activated
blowing agent.

The chemical embossing inhibitors embodied in this
invention have the advantage that they are insoluble or
substantially insoluble in water, water/alcohol mixtures

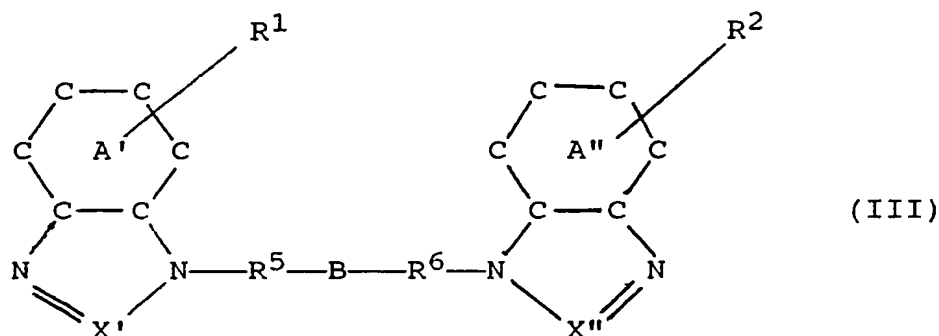
and many organic solvents, and can be used to form stable dispersions which do not adversely effect the stability and printing characteristics of either anionic or cationic aqueous printing inks of widely varying compositions. Ink compositions according to the invention preferably comprise water as the solvent. The compounds are also significantly less prone to uncontrolled lateral migration and migration through the foamable substrate than the commonly used benzo- and tolyltriazole inhibitors. Therefore, the resulting image is sharper and more distinct, as well as ghost embossing being reduced.

The preferred structures of the highly insoluble azoles of this invention are those according to the formula (I) in which the A ring is a benzenoid, R is a methyl group, R¹ is a hydrogen or methyl and X is a nitrogen atom. The most active inhibitors of the present invention which have been made are those having a 1-methyl benzotriazole moiety attached to a nitrogen atom and a second 1-methyl benzotriazole, carboxy containing or sulfonyl linking moiety attached to the same or different nitrogen atom.

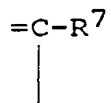
Advantageously, the structures of the highly insoluble azoles of this invention are those according to formula (I), especially as described directly above, wherein Y is a hydrogen atom, an alkyl group of 1 to 4 carbon atoms, an hydroxyl moiety or a carboxy containing moiety and Z is methyl benzotriazole, methyl

tolyltriazole, a carbonyl or thiocarbonyl linked methyl benzotriazole or methyl tolyltriazole containing moiety or a sulfonyl linked moiety or Y and Z taken together form a saturated ring compound containing a carbonyl group, or having a methyl benzotriazole or methyl tolyltriazole containing moiety attached thereto.

Advantageously, ink compositions according to the invention comprise an inhibitor having the formula



wherein each of the A' and A'' rings, which may be the same or different, is a benzenoid, naphthenoid or saturated cycloaliphatic moiety and are each unsubstituted or substituted with R¹ or R², which may be the same or different, and each represents an alkylene group of 1 to 4 carbon atoms; R⁵ and R⁶, which may be the same or different, each represents an alkylene group of 1 to 5 carbon atoms, X' and X'', which may be the same or different, each represents a nitrogen atom or the



group wherein R⁷ is a hydrogen atom or an alkyl group of 1 to 4 carbon atoms; B is NH, NOH or an organic moiety; and both the R⁵-B and B-R⁶ bonds are carbon/nitrogen

bonds. X' is preferably a nitrogen atom and more preferably X' and X'' are both nitrogen atoms. Preferably A' and A'' are the same and R¹ and R² are the same. R⁵ and R⁶ are preferably the same.

Preferably B in formula (III) represents organic substituted nitrogen, organic substituted nitrogen/carbonyl/nitrogen, organic substituted nitrogen/carbonyl/nitrogen/carbonyl/nitrogen, or an organic substituted or unsubstituted saturated heterocyclic ring.

Ink compositions according to the present invention preferably comprise an inhibitor which is a compound having

- (a) at least two moieties selected from the group consisting of 1-methyl benzotriazole moiety, 1-ethyl benzotriazole moiety, 1-methyl benzimidazole moiety, 1-ethyl benzimidazole moiety, 1-methyl tolyltriazole moiety and 1-ethyl tolyltriazole moiety, the selected moieties being attached to one or more nitrogen atoms; or
- (b) at least one moiety selected from the group consisting of 1-methyl benzotriazole moiety, 1-ethyl benzotriazole moiety, 1-methyl tolyltriazole moiety and 1-ethyl tolyltriazole moiety, the selected moiety being attached to a nitrogen atom, and the inhibitor being a compound having at least one carboxy moiety; or

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(c) at least one moiety selected from the group consisting of 1-methyl benzotriazole moiety, 1-ethyl benzotriazole moiety, 1-methyl tolyltriazole moiety and 1-ethyl tolyltriazole moiety, the selected moiety being attached to a nitrogen atom; and the inhibitor being a compound having at least one sulfonyl moiety attached to a nitrogen atom.

The inhibitors which have been tested and found to have inhibitor activity include 1,3-bis(benzotriazol-1'-yl methyl) urea; 1,3-bis(5'-tolyltriazol-1'-yl methyl) urea; 1,5-bis(benzotriazol-1'-yl methyl) biuret; 2,4,6-

tris(benzotriazol-1'-yl methyl)-s-triazine; 2,4-
 bis(benzotriazol-1'-yl methyl) benzoguanamine; 1,3-
 bis(benzotriazol-1'-yl methyl) N,N'-dimethyl urea; 1-(1'-
 methanesulfonamido)methyl benzotriazole; 1-(1'-
 5 benzenesulfonamido) methyl benzotriazole; 4-(benzotriazol-1'-
 yl methyl) hydantoin; 1-(1'-(2'-oxopyrrolidin-1'-yl) ethyl)
 benzotriazole; N,N-bis(benzotriazol-1-yl methyl)
 hydroxylamine; N-(benzotriazol-1-yl methyl) 4'-carboxybenzene
 sulfonamide; N,N-bis(benzotriazol-1-yl methyl) glycine; 1,3-
 10 bis(benzotriazol-1'-yl methyl) thiourea; N,N'-bis(benzotriazol-1-yl
 methyl) naphthalene-1,5-disulfonamide; N,N'-bis(tolyltriazol-1-
 yl methyl) piperazine; N,N'-bis(benzotriazol-1-yl methyl)
 piperazine; N,N'-bis(methylcyclohexyltriazol-1-yl methyl)
 piperazine; N,N'-bis(benzotriazol-1-yl methyl) benzene-1,3-disulfonamide;
 15 1-bis(benzotriazol-1'-yl methyl)-2-benzoyl
 hydrazide; bis(benzotriazol-1-yl methyl) amine; 1,3-bis(5'-
 butyl benzotriazol-1'-yl methyl) urea; benzimidazole; and
 N,N'-bis(benzimidazol-1-yl methyl) piperazine. These inhibitors are
 preferred. The following compounds are new compounds, not known to the
 20 present inventors to have been previously synthesized, 1,3-bis(5'-
 tolyltriazol-1'-yl methyl) urea; 1,5-bis(benzotriazol-1'-yl
 methyl) biuret; 2,4,6-tris(benzotriazol-1'-yl methyl)-s-
 triazine; 2,4-bis(benzotriazol-1'-yl methyl) benzoguanamine;
 N,N-bis(benzotriazol-1-yl methyl) glycine; N,N'-bis(benzotriazol-1-yl
 25 methyl) naphthalene-1,5-disulfonamide; N,N'-bis(benzotriazol-1-
 yl methyl) piperazine; N,N'-bis(methylcyclohexyltriazol-1-yl
 methyl)piperazine; N,N'-bis(benzotriazol-1-yl methyl) benzene-1,3-

disulfonamide; 1-bis(benzotriazol-1'-yl methyl)-2-benzoyl
hydrazide; bis(benzotriazol-1-yl methyl) amine; 1,3-bis(5'-
butyl benzotriazol-1'-yl methyl) urea; N,N'-bis(benzimidazol-
1-yl methyl) piperazine; and N-(benzotriazol-1-yl methyl)-4'-
5 carboxybenzene sulfonamide. It is expected
that N,N'-bis(cyclohexyltriazol-1-yl methyl) piperazine would
be an effective inhibitor. These inhibitors are preferred.

For acceptable processing, it is advantageous to use 1 to
15 percent by weight of the insoluble azole dispersed in the
10 aqueous printing ink composition, and preferably 5 to 10
percent by weight, especially for floor covering applications. Higher
concentrations can be used (>15%) depending on the application
weight of the wet applied ink. Shallower engraved cylinders
may require more inhibitor per unit area to get the desired
15 embossed effect.

Those skilled in the art will recognize that a very wide
range of printing ink compositions exist with varying
combinations of resin binders, pigments, inhibitors and
rheology-control additives. The pigments are optional, since
20 it may be desirable to use a colorless, inhibitor containing,
printing ink. The insoluble azole compounds of this invention
are potentially useful in many other aqueous or solvent ink
formulations not specifically outlined in the Examples as to
their exact composition.

25 Those skilled in the art will also recognize that varying
amounts of water will be required to adjust the viscosity of
the ink composition to a range suitable for typical

rotogravure printing. Other methods of printing the ink composition onto the foamable plastic surface, such as screen printing, relief printing, or planographic printing, may also be used with these ink compositions.

5 Although this invention is primarily concerned with polyvinylchloride-based plastisol compositions thermally blown with azodicarbonamide or other blowing agents as the printing substrate, there likewise exists a wide range of resins which can be thermally foamed with azodicarbonamide and thus are
10 potential substrates for aqueous inhibitor printing ink compositions of the type claimed. Such other compositions include polyvinyl acetate, copolymers of vinyl chloride and vinyl acetate, polyacrylate, polymethacrylate, polyethylene, polystyrene, butadiene/styrene copolymers,
15 butadiene/acrylonitrile copolymers, and natural or synthetic rubbers.

 The specific combinations of PVC, other resins, filler, stabilizers, plasticizers, chemical blowing agents and activators that make up a typical foamable plastisol substrate
20 vary widely within certain limits.

 The invention is illustrated by the following examples related to synthesis of the insoluble azole derivatives,
25 preparation of the aqueous dispersions and printing ink formulations, and demonstration of the chemical embossing behavior of the claimed compounds. Unless otherwise stated,

all amounts and percentages given in the Examples are on a weight basis.

EXAMPLE 1

Preparation of N,N'-Bis(Tolyltriazol-1-yl Methyl) Piperazine (TTA-P)

In a flask, were combined 133.13 parts of commercial tolyltriazole (TT100, an isomer mixture from PMC Specialties) and 43.1 parts piperazine in 700 parts methanol and cooled to zero degrees Centigrade. While holding the reaction mixture at this temperature, 81.2 parts of commercial aqueous 37% formaldehyde solution was added slowly over several hours with continual agitation, during which time a finely divided white solid began to precipitate. The reaction mixture was allowed to warm to ambient temperature and worked up after 48 hours by suction filtration. The filter cake was washed once by suspending the solid in a fresh charge of methanol and applying vacuum to remove the liquid. The resulting material was dried under moderate vacuum at 65-75°C to give 181.7 parts (96.5% yield) of a white powdery solid which was identified by standard spectroscopic techniques as TTA-P; N,N'-bis(tolyltriazol-1-yl methyl) piperazine.

EXAMPLE 2

Preparation of N,N'-Bis(Benzotriazol-1-yl Methyl) Piperazine (BTA-P)

In a flask, were combined 119.13 parts of commercial benzotriazole (Cobratec 99 from PMC Specialties) and 43.1 parts piperazine in 500 parts methanol and treated with 81.2 parts of commercial aqueous 37% formaldehyde solution at zero

degrees Centigrade as in Example 1. After 48 hours at room temperature, the resulting solid product was filtered, methanol washed and dried under moderate vacuum at 65-75°C to give 170.4 parts (97.8% yield) of white powder which was identified by standard spectroscopic techniques as BTA-P; N,N'-bis(benzotriazol-1-yl methyl) piperazine.

EXAMPLE 3

Preparation of N,N'-Bis(Methylcyclohexyltriazol-1-yl Methyl) Piperazine (HTTA-P)

In a flask, were combined 178.1 parts of hydrogenated tolyltriazole (Cobratec 911 from PMC Specialties) and 55.1 parts piperazine in 500 parts methanol and treated with 103.8 parts of commercial aqueous 37% formaldehyde solution at zero degrees Centigrade as in Examples 1 and 2. After 48 hours at room temperature, the resulting solid product was filtered, methanol washed and dried under moderate vacuum at 65-75°C to give 200.1 parts (80.5% yield) of white powder which was identified by standard spectroscopic techniques as HTTA-P; N,N'-bis(methylcyclohexyltriazol-1-yl methyl) piperazine.

EXAMPLE 4

Preparation of Cationic Dispersion of TTA-P

A cationic dispersion of TTA-P was prepared using a quaternary ammonium salt, stearyl dimethylbenzylammonium chloride, (Maquat SC-18, Mason Chemical Co.), as the stabilizer. The product was first diminutized by grinding the coarse powder (TTA-P) for approximately 18 hours in a standard ball mill using a combination of 12 mm diameter spherical and

6 mm diameter X 6 mm high cylindrical balls. Approximately 1/2 of the 1 L ball mill volume was filled for the grinding operation. After milling, microscopic observation showed reduction of particle size from 30-50 microns to 1-10 microns.

5 The dispersion was then prepared by adding 2.35 parts Maquat SC-18 (85% active) to 37.65 parts deionized water and stirring until dissolved. A total of 40 parts of TTA-P was then added to the surfactant solution in 5 part increments with stirring, followed by agitation with a sonic dismembrator, (Fisher Model

10 3000). The sonic probe was inserted directly into the suspension and run on the highest setting for 1-2 minutes. A creamy dispersion resulted initially, with viscosity increasing with increasing solids content. The final suspension was a uniform paste with a concentration of 50% by

15 weight solids.

EXAMPLE 5

Preparation of Inhibited Cationic Aqueous Rotogravure Ink Formulation with TTA-P

A blue aqueous inhibitor ink was prepared by adding 0.20

20 parts of CIB 103 Blue Pigment (sold by Penn Color, Inc.) to 20 parts of CIE 94 Extender (sold by Penn Color, Inc.) and stirring to uniform coloration. Then 6.06 parts of the 50% suspension of TTA-P (prepared in Example 4) was then added to the ink mixture and stirred to uniform coloration. Although a

25 slight viscosity drop was observed the mixture remained colloiddally stable and disperse.

EXAMPLE 6

Preparation of an Anionic Dispersion of TTA-P

An anionic dispersion paste of TTA-P was prepared with a
5 polyoxyethylene branched nonylphenyl ether phosphate
surfactant, (Rhodofac PE-50, Rhone Poulenc). Diminutization
of the compound was performed by milling as described in
Example 4. Dispersion of the compound was accomplished using
the sonic dismembrator, also described in Example 4. Materials
10 were combined in the proportion: 40 parts of deionized water,
2.06 parts of the surfactant and 40 parts of TTA-P. Materials
were added in the listed sequence. A stable, homogenous
paste, 48.7% TTA-P, resulted. Microscopic observation showed
particles approximately 1-3 microns in diameter.

EXAMPLE 7

Preparation of an Inhibited Anionic Aqueous Rotogravure Ink Formulation with TTA-P

The anionic dispersion of TTA-P (Example 6) was added to
an anionic ink formulation of Sicpa Corp. The resultant
20 mixture consisted of 20 parts Sicpa Extender 694556, 0.20
parts Sicpa black ink 674554 and 6.06 parts of the 48.7%
dispersion of the compound (Example 6). The mixture was
stirred to a smooth and uniform consistency, and was observed
to be colloidally stable.

EXAMPLE 8

Direct Addition of TTA-P to an Aqueous Cationic Ink Formulation

Hitherto, dispersion of these new inhibitors directly
5 into the ink formulation was not considered. The instability
problems of other triazole inhibitors seemed to indicate that
a surfactant in addition to that found in the ink and extender
was required. But since such stable ink mixtures were
obtained with addition of the charged dispersions it was
10 thought to attempt the dispersion of a new triazole inhibitor
directly into an ink mixture. The attempt was successful. To
20 parts of Extender CIE 94 were added 0.20 parts of CIB 103
blue ink and the mixture was stirred to a smooth uniform
coloration. Three parts of the milled tolyltriazole-
15 piperazine derivative, TTA-P (milled as in Example 4) were added
directly to the ink mixture and the mixture was sonicated to a smooth
consistency. Sonication was performed discontinuously to
avoid overheating and coalescence of extender latexes. A
homogeneous and stable mixture was obtained.

EXAMPLE 9

Direct Addition TTA-P to an Aqueous Anionic Ink Formulation

To 20 parts of Sicpa extender 694556 was added 0.20 parts
of Sicpa 674554 black ink and the mixture was stirred to a
25 uniform coloration. Three parts of TTA-P (milled as in
Example 4) were added and the mixture was sonicated to a
smooth consistency. A homogenous and stable mixture was
obtained.

EXAMPLES 10 - 13

Printing of Inks and Resultant Embossing

The inks prepared in Examples 5, 7, 8 and 9 were printed
5 on 9 mils (0.23 mm) of an expandable plastisol coated on flooring felt
using a flat-bed gravure proof press. The plastisol
formulation was 100 parts PVC resin, 50 parts plasticizers, 30
parts limestone filler, 7.0 parts titanium dioxide pigment,
3.0 parts mineral spirits viscosity modifier, 2.1 parts
10 stabilizers, 2.0 parts azodicarbonamide blowing agent and 0.6
parts zinc oxide blowing agent activator. The inks printed
and dried satisfactorily without any tack.

The printed samples were coated with 10 mils (0.25 mm) of a clear
plastisol and heated for 1.3 ± 0.1 minutes at an air
15 temperature of $201 \pm 1^\circ\text{C}$ in a Werner Mathis oven to expand the
9 mil (0.23 mm) layer to about 25 mils (0.64 mm). The clear plastisol
formulation was 100 parts PVC resin, 40 parts plasticizers, 4.0 parts
stabilizers and 4.0 parts mineral spirits viscosity modifier.

The thickness of the printed coated areas (i.e.,
20 restricted area) was measured in mils (mm) and compared to the
thickness of the unprinted expanded surrounding areas. This
difference is reported as the depth of chemical embossing and
is shown in Table I.

TABLE I

		<u>Ink</u>	<u>Weight Percent of Compound in Ink</u>	<u>Chemical Embossing Depth in mils (mm)</u>	
<u>EXAMPLE</u>					
5	10	Cationic Ink (Example 5)	11.54 TTA-P	10.0	(0.254)
	11	Anionic Ink (Example 7)	11.24 TTA-P	9.1	(0.231)
	12	Cationic Ink (Example 12)	12.93 TTA-P	10.6	(0.270)
10	13	Anionic Ink (Example 13)	12.93 TTA-P	10.8	(0.274)

EXAMPLE 14

Direct Milling of BTA-P
In an Aqueous Cationic Ink Formulation

15 The following procedure was developed to see if the
present inhibitors could be directly milled into water-based
ink systems. Instead of pregrinding and then dispersing the
inhibitor as was done in Example 4, the compound from Example
2 (BTA-P) was ground and dispersed in situ in the water-based
20 ink extender. A sixteen ounce (454 g) HDPE bottle was filled halfway
with a mixture of 12 mm diameter spherical and 6 mm diameter X
6 mm high cylindrical ceramic balls. To the bottle was added
21.6 grams of the coarse powder of BTA-P and then 158.4 grams
of extender CIE 94 from Penn Color, Inc. This gave a
25 concentration of 12% by weight of BTA-P and room to adjust the
concentration and viscosity with water and more extender.

The charged mill was rolled overnight (about 18 hours)
and checked for the quality of the grind. A homogeneous
stable dispersion was obtained and under microscopic

observation showed particle size reduction from over 50 microns to less than 10 microns. The ceramic balls were separated from the dispersion and the dispersion adjusted to about 10% by weight concentration of BTA-P with water and additional extender to a viscosity of 15 seconds with a #3 Zahn Cup. The morphological properties of the compound and lack of solubility in the ink allow the material to be readily ground and dispersed in situ.

10	<u>Formulation Before Milling</u>	<u>Formulation After Milling and Adjusting Viscosity</u>
	88 parts Extender CIE 94	79.parts Extender CIE 94
	12 parts BTA-P	9.6 parts BTA-P
		10.6 parts Water

EXAMPLE 15

15 Preparation of 1,3-Bis(Benzotriazol-1'-yl Methyl) Urea (BTA-U)

In a flask, were combined with stirring 119 parts of benzotriazole and 30 parts of urea in a solution of 150 parts of water and 200 parts of glacial acetic acid at room temperature. To the resulting clear, pale yellow solution that had cooled to about 15°C, from dissolution of urea and benzotriazole, was added in about one hour 89 parts of aqueous 37% formaldehyde. Approximately 2/3 through the addition of formaldehyde, a finely divided white solid began to precipitate. Upon completing the addition, the reaction temperature had risen to 35°C. Stirring was continued for several hours.

After about 16 hours at room temperature, the reaction mixture was suction filtered. The white solid filter-cake was washed consecutively with portions of a 50/50 (by vol.) aqueous/acetic acid solution and finally water. Air drying of the washed filter-cake, followed by drying in vacuo (in presence of phosphorus pentoxide) provided 126 parts (78% yield) of a white solid, m.p. 221-223°C. The material was identified by ¹H and ¹³C NMR spectral analysis as 1,3-bis(benzotriazol-1'-yl methyl) urea.

EXAMPLE 16

Preparation of 1,3-Bis(5'-Tolyltriazol-1'-yl Methyl) Urea (5-TTA-U)

The previous reaction was repeated using 53.5 parts of 5-tolyltriazole and 10.5 parts of urea in 70 parts of acetic acid and 55 parts of water. To this stirred mixture was added 32.4 parts of aqueous 37% formaldehyde. The resulting reaction mixture was subsequently heated to 60°C and maintained at this temperature for about 18 hours. The reaction mixture was allowed to cool to room temperature and washed consecutively with water, methanol and ether. After drying in vacuo, the product, 60 parts, melted at 184-8°C, and was identified as 1,3-bis(5'-tolyltriazol-1'-yl methyl) urea (98% yield) by ¹H and ¹³C NMR spectral analysis.

EXAMPLE 17

Preparation of 1,3-Bis(Benzotriazol-1'-yl Methyl) N,N'-Dimethyl Urea (BTA-DMU)

To a solution of dry toluene (250 parts) and p-toluene
5 sulfonic acid (1.7 parts) was added 8.8 parts of dimethyl urea
and 59.6 parts of 1-(hydroxymethyl) benzotriazole. The
stirred mixture was heated to reflux under a Dean-Stark trap
and became clear. Refluxing was continued for 24 hours, after
which time the reaction mixture was cooled to room
10 temperature. The reaction mixture was washed consecutively
with portions (50 parts) of aqueous 5% sodium carbonate, water
and aqueous saturated sodium chloride and finally dried over
anhydrous magnesium sulfate. The dried and filtered solution
was concentrated at reduced pressure to provide a viscous oil.
15 1,3-Bis(benzotriazol-1'-yl methyl) N,N'-dimethyl urea was
isolated from the oil, m.p. 137-40°C (reported m.p. 143-4°C).
NMR spectral analysis of the product corresponded with that
reported in the literature.

EXAMPLE 18

20 Preparation of 2,4,6-Tris(Benzotriazol-1'-yl Methyl)-s-Triazine (3BTA-M)

To a stirred mixture of melamine (37.8 parts) and 107.2
parts of benzotriazole in acetic acid (315 parts) and water
(225 parts) was added in 20 minutes aqueous 37% formaldehyde
25 (74.2 parts). Upon completing the addition, the stirred
reaction mixture was heated to 45°C and maintained for 19
hours. The reaction mixture was cooled and filtered with
suction. The filter-cake was washed consecutively with water,

methanol and ether and dried in vacuo at 55°C. The dried product, m.p. 226-30°C, 130 parts (83.4%) was identified as 2,4,6-tris(benzotriazol-1-yl methyl)-s-triazine by ¹H and ¹³C NMR spectral analysis.

5

EXAMPLE 19

Preparation of 1-(1'-Benzenesulfonamido) Methyl Benzotriazole (BTA-BSA)

A mixture of benzenesulfonamide (47.2 parts) and 1-hydroxymethylbenzotriazole (46.2 parts) in 400 parts of dry
10 toluene was refluxed under a Dean-Stark trap. After about 24 hours, a near theoretical amount (5.1 parts) of water had formed. The reaction mixture was cooled to room temperature. A white solid present was filtered, washed with fresh toluene and dried in vacuo to provide 80.6 parts (93.2 % of theory) of
15 1-(1'-benzenesulfonamido) methyl benzotriazole. The product melted at 180-3°C (reported m.p. 183-6°C) and was further characterized by ¹H and ¹³C NMR.

EXAMPLE 20

Preparation of N,N-Bis(Benzotriazol-1-yl 20 Methyl) Hydroxylamine (BTA-NOH)

To a stirred solution of 1-(hydroxymethyl)benzotriazole (44.8 parts) in 375 parts of methanol at room temperature was added 10.4 parts of hydroxylamine hydrochloride. The reaction mixture was stirred at room temperature for about five hours
25 and then placed in a freezer for about six hours. The precipitated white solid was filtered, washed with cold water and dried in vacuo (in presence of phosphorus pentoxide). The dried product, m.p. 175-7°C (reported m.p. 173-4°C), 24 parts,

was identified as N,N-bis (benzotriazol-1-yl methyl) hydroxylamine (54.2% yield) by ¹H and ¹³C NMR spectral analysis.

EXAMPLE 21

5 Preparation of 1,3-Bis(Benzotriazol-1'-yl Methyl) Thiourea (BTA-TU)

To a stirred mixture of benzotriazole (119 parts) and thiourea (38 parts) in 300 parts of acetic acid at room temperature was added 89 parts of aqueous 37% formaldehyde in
10 about one hour. Upon completing the addition, the reaction mixture was heated to about 55°C. After 12 hours at 55°C, the reaction mixture was cooled to room temperature and the solid present was suction filtered. The filter-cake was washed consecutively with water, methanol and ether. The solid
15 product was dried in vacuo to provide 161 parts (95% yield) of 1,3-bis(benzotriazol-1'-yl methyl) thiourea, m.p. 220-2°C (reported m.p. 205-6°C), and identified by ¹H and ¹³C NMR spectral analysis.

EXAMPLE 22

20 Preparation of N,N-Bis(Benzotriazol-1-yl Methyl) Glycine (BTA-G)

One hundred and nineteen and three tenths (119.3) parts of 1-(Hydroxymethyl) benzotriazole and glycine (30 parts) were added to 600 parts of dry toluene containing 1.7 parts of
25 p-toluenesulfonic acid. The mixture was stirred and refluxed under a Dean-Stark trap. After about 4.5 hours, the theoretical amount of water (14.4 parts) had collected and heating was suspended. The reaction mixture was cooled in

ice-water and the tan solid that had formed was filtered with suction. After washing the filter-cake consecutively with toluene and ether and drying it in vacuo, 116 parts of N,N-bis(benzotriazol-1-yl methyl) glycine were obtained and identified by ¹H and ¹³C NMR spectral analysis. The product melted at 163-6°C and was obtained in 86% yield.

EXAMPLE 23

Preparation of N-(Benzotriazol-1-yl Methyl)- 4'-Carboxybenzene Sulfonamide (BTA-4CBSA)

To a stirred mixture of benzotriazole (23.8 parts) and 4-carboxybenzene sulfonamide (40.2 parts) in acetic acid (250 parts) was added 17 parts of aqueous 37% formaldehyde in 25 minutes. The resulting reaction mixture was heated to 55°C. After about 18 hours at 55°C, the reaction mixture was cooled to room temperature and the white solid present was filtered with suction. The filter-cake was washed consecutively with portions of water, methanol and ether. After drying in vacuo, 60 parts of a solid melting at 258-61°C was obtained. The solid was identified as N-(benzotriazol-1-yl methyl)-4'-carboxybenzene sulfonamide (90% of theory) by ¹H and ¹³C NMR spectral analysis.

EXAMPLE 24

Preparation of 1(1',5'-Naphthalene Disulfonamido) Methyl Benzotriazole (BTA-NDSA)

[N,N'-bis(benzotriazol-1-yl-methyl) naphthalene-1,5-disulfonamide]

To a solution of dry toluene (400 parts) and p-toluene sulfonic acid (0.5 parts) was added 42.9 parts of 1,5-naphthalene disulfonamide and 46.2 parts of 1-(hydroxymethyl) benzotriazole. The stirred reaction mixture was heated under

reflux under a Dean-Stark trap and refluxing was continued for 8 hours. The reaction mixture was cooled, filtered and washed with cold methanol. Attempts to recrystallize the material were unsuccessful due to the insolubility of the product in many organic solvents (hot and cold). The material was heated in methanol and filtered hot. The white solid was dried in a vacuum oven and yielded 58.5 parts of a material with a melting range of 245-50°C with darkening. The material was identified as 1(1',5'-naphthalene disulfonamido) methyl benzotriazole by ¹H and ¹³C NMR spectral analysis, run in DMSO-d₆.

Example 25

Preparation of 1(1',3'-Benzene Disulfonamido) Methyl Benzotriazole (BTA-BDSA)

[N,N'-bis(benzotriazol-1-yl-methyl) benzene-1,3-disulfonamide]

To a solution of dry toluene (400 parts) and p-toluene sulfonic acid (0.5 parts) was added 38.7 parts of 1,3-benzene disulfonamide and 50.5 parts of 1-(hydroxymethyl) benzotriazole. The stirred reaction mixture was heated under reflux under a Dean-Stark trap and refluxing was continued for 8 hours. The reaction mixture was cooled, filtered and washed in boiling methanol. Attempts to recrystallize the material were unsuccessful due to the insolubility of the product in many organic solvents. The precipitate was dried in a vacuum oven to yield 41.8 parts of a material which began to darken at 240°C and melted in the range of 255-60° with gas evolution. The material was identified as 1(1',3'-benzene

disulfonamido) methyl benzotriazole by ^1H and ^{13}C NMR spectral analysis, run in DMSO- d_6 .

EXAMPLE 26

Preparation of 1-Bis(Benzotriazol-1'-yl Methyl)-2-Benzoyl Hydrazide (BTA-HYR)

A mixture of benzoic hydrazide (42.7 parts) and 1-(hydroxymethyl)benzotriazole (104.4 parts) in 600 parts of dry benzene was heated to reflux with stirring. After about 4 hours, approximately 80% of the theoretical amount of water had formed and the heating was terminated. Upon cooling, a white solid that had formed was filtered with suction, washed consecutively with portions of methanol and ethyl ether and finally dried in vacuo. The dried product, 132.4 parts, m.p. 217-220°C was identified as 1-bis(benzotriazol-1'-yl methyl)-2-benzoyl hydrazide (95% yield)) by ^1H and ^{13}C NMR spectral analysis.

EXAMPLE 27

Preparation of Bis(Benzotriazol-1-yl Methyl) Amine (BTA-A)

An aqueous 2% ammonia solution (265 parts) was neutralized with acetic acid using phenolphthalein indicator. To the resulting solution at 25°C was added a solution of 1-(hydroxymethyl)benzotriazole (74.6 parts) in about 600 parts of methanol. The reaction mixture was stirred at 25°C for 5 hours and then placed in a freezer (-5°C) overnight.

The solid precipitate that formed was filtered, washed with ice water and dried in vacuo in the presence of phosphorus pentoxide. The dried white solid, 15.8 parts,

melted at 182-5°C. The main methanol filtrate was concentrated to 1/2 its original volume and then cooled in the freezer. The ice cold concentrate was filtered and the filter-cake treated as described above provided 18.6 parts of a white solid, m.p. 182-5°C. The combined solids, 34.4 parts, was identified as bis(benzotriazol-1-yl methyl) amine (49% of theory) by ¹H and ¹³C NMR spectral analysis.

EXAMPLE 28

Preparation of 1,3-Bis(5'-Butyl Benzotriazol-1'-yl Methyl) Urea (5-BBTA-U)

Urea (8.6 parts) and 5-butyl benzotriazole (50.0 parts) were added to 100 parts of glacial acetic acid. Aqueous 37% formaldehyde (23.1 parts) was added dropwise and upon completion of addition, the mixture was heated to 60°C and stirred overnight at this temperature. The system was cooled and the precipitate was suction filtered, washed with water and dried in a vacuum oven to yield 36.2 parts of solid melting at 157-161°C.

EXAMPLE 29

Preparation of N,N'-Bis(Benzimidazol-1-yl Methyl) Piperazine (BI-P)

Benzimidazole (121.7 parts) and 43.1 parts of piperazine were mixed in 600 parts of methanol and cooled to 0°C. While holding the reaction mixture at 0 to 12°C, 81.2 parts of commercial aqueous 37% formaldehyde solution were added over several hours with continuous stirring. After addition, the system was allowed to warm to room temperature. The system was allowed to stand overnight and then suction filtered. The

solid was washed with methanol and placed in a vacuum oven to yield 162.0 parts of material melting at 250-3°C.

Table II sets forth a number of properties of the examples and comparative examples which were made and tested.

5 Most of these compounds were prepared using the direct milling procedure and then evaluated for inhibitor activity, ghosting and embossing definition.

The inhibited inks (at 10% by weight inhibitor concentration) were printed on 7 mils (0.18 mm) of an expansible
10 plastisol coated onto a glass mat which was saturated with a non-expandsible plastisol. This was done on a flat-bed gravure proof press using a 100 line screen step-wedge gravure plate. The steps ranged from a deep shadow tone to a shallow highlight tone. The inks printed and dried without any tack.

15 The printed samples were coated with 10 mils (0.25 mm) of a clear plastisol wearlayer and heated for 1.9 ± 0.1 minutes at an air temperature of $185 \pm 2^\circ\text{C}$ in a Werner Mathis oven to fuse and expand the foamable plastisol to about 14 mils (0.36 mm) (a 2:1 blow ratio). The thickness of the printed coated areas (i.e.,
20 restricted area) was measured in mils (mm) and compared to the thickness of the expanded unprinted surrounding areas. This difference was recorded as depth of chemical embossing and was used along with the degree of expansion in the inhibited area to assess the inhibitor activity (IA).

25 The inhibitor activity of the BTA-P derivative was established as the benchmark and on a scale of 1 to 5 was given a rating of 1 (five on the scale being less than one mil

(0.025 mm) of overall chemical embossing). This is a subjective ranking where the other compounds were evaluated for inhibitor activity by comparing them to BTA-P, both numerically and visually.

5 Those compounds showing good inhibitor activity were also evaluated for nonghosting characteristics. Ghosting is a result of "in roll" migration of the inhibitor from one printed surface of a rolled sheet into the lap above. Migration and ghosting also occurs in the other direction
10 (i.e., to the lap below) but not as rapidly. The result of this fugitive migration is an embossed image (ghost embossing) showing up after expansion in an area not printed with the inhibited ink. This phenomenon is readily seen with
15 inhibitors like benzotriazole and tolyltriazole in rolls of printed flooring structures after a few hours or days. Structures that have vinyl plastisol throughout are more prone to this problem.

To speed up the evaluation of ghosting, a bench top test was developed. Printed samples were held under pressure at an
20 elevated temperature of 120°F (49°C) for the desired period of time at 1.4 psi (984 kg/m²). A multi-layer sample stack was compressed between two 3/4 inch (19 mm) plywood boards to distribute the pressure uniformly. This simulated the conditions rolls of printed material could be stored under before expansion. The elevated
25 temperature accelerated the migration and showed results in hours or days rather than days or weeks at room temperature.

Testing consisted of printing the inhibited inks on the flooring structure, as described previously with respect to inhibitor activity, using a grout line engraved plate on a flat-bed gravure proof press. Printed samples were sandwiched between unprinted sheets of the same flooring structure and placed in an air-circulating oven under heat and pressure. Unprinted sheets were used to make it easier to see ghosting when it first started to occur.

The samples were removed from the oven over a period of time (e.g., hours, days or weeks) and expanded in a Werner Mathis oven at $185 \pm 2^{\circ}\text{C}$ for 1.9 ± 0.1 minutes. The top and bottom unprinted sheets were evaluated for signs of ghosting. When ghosting occurred, a slight to severe embossed image of the grout line could be seen.

SL = Slight, faint discontinuous print image with very little embossing.

M = Moderate, faint continuous print image with little embossing.

S = Severe, ghost embossing nearly equal to the direct printed samples.

In addition to evaluating ghosting over time, embossing definition can also be evaluated using the printed sheets from the ghosting test. The printed sheets were expanded at the same time intervals as the unprinted ghosting sheets and evaluated for depth and sharpness of the printed/embossed image. It was found that those inhibitors with severe ghosting characteristics (e.g. BTA and TTA) showed poor

embossing definition over time. This is attributed to the lateral migration of the inhibitor and the depletion of the inhibitor in the print area.

TABLE II

5	<u>EXAMPLE</u>	<u>IA^a</u>	<u>MW</u>	<u>MP°C</u>	<u>GHOSTING^b</u>	<u>SOLUBILITY</u> <u>% BY WT^c</u>	
						<u>H₂O</u>	<u>iPrOH</u>
	TTA-P	1	376.4	194-7	N	0.031	0.204
	BTA-P	1	348.4	>215	N	0.001	0.007
10	HTTA-P	1	388.4	145	N	-	-
	BTA-U	<1	322.3	221-3	N	0.002	0.08
	5-TTA-U	3	350.3	184-8	-	-	-
	BTA-DMU	3	350.4	137-40	SL	-	-
	3BTA-M	3	519.5	226-30	-	-	-
15	BTA-BSA	>1	289.3	180-3	S	0.05	0.59
	BTA-NOH	<1	295.3	175-7	S	0.08	1.08
	BTA-TU	2	338.3	220-2	N	0.001	0.011
	BTA-G	>1	337.4	167-9	S	0.69	0.914
	BTA-4CB SA	>1	332.3	258-61	SL	0.011	0.137
20	BTA-NDSA	1	548.0	245-50	N	0.051	0.051
	BTA-BDSA	>1	498.0	240-50	N	0.089	0.120
	BTA-HYR	1	398.4	217-20	N	0.006	0.019
	BTA-A	>1	279.3	182-5	S	0.039	0.028
	5-BBTA-U	3	434.4	157-61	-	-	-
25	BI ^d	2	118.1	172-4	S	0.50	15.30
	BI-P	1	346.0	250-3	N	0.053	0.211
	Comparative Examples						
	BTA ^e	2	119.2	98-9	S	1.98	53.9
	TTA ^f	2	133.2	83-5	S	0.55	52.9
30	TTA-HE ^g	3	250.2	52-4	S	abt 50	>50
	TTA-EH ^h	2	386.2	<25	S	<0.01	>50

^a IA-Inhibitor Activity - 1=Excellent, 2=Very Good, 3=Good,
4=Fair, 5=Poor and N=None

^b N=None, SL=Slight and S=Severe (after 3 days)

^c At room temperature for 24 hours

5 ^d Benzimidazole

^e Benzotriazole

^f Tolyltriazole

^g 1-Bis(β-hydroxyethyl)aminomethyltolyltriazole, (Reomet 42,
trademark of Ciba-Geigy)

10 ^h 1-Bis(2-ethylhexyl)aminomethyltolyltriazole, (Reomet 39,
trademark of Ciba-Geigy) (Liquid at room temp.)

Due to the extremely low solubilities of the present
compounds in both water and alcohol, and the fact that they
are solid particles at room temperature, they can be treated
15 like pigments in any ink composition. The present compounds
do not lead to instability of the ink and may be dispersed
into the ink composition by either micronizing and dispersing
or simultaneously grinding and dispersing.

The compounds of the prior art are either liquids at room
20 temperature or are sufficiently soluble in water or alcohol to
make it infeasible to mix the prior art inhibitors into the
ink composition and then simultaneously grind and disperse
them in the ink composition. Therefore, the presently claimed
insoluble azole inhibitors have a major commercial advantage
25 over the prior art inhibitors.

While only some of the imidazole compounds have been
tested, it is believed that the imidazole compounds
corresponding to the triazole compounds would be effective
insoluble inhibitors. However, the triazole compounds are

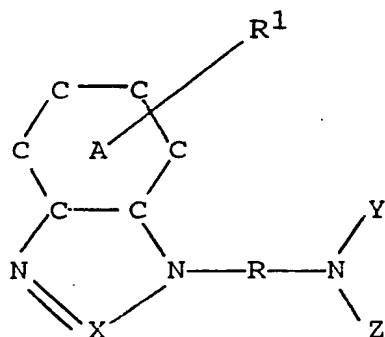
preferred since at least some of the corresponding imidazole compounds appear to lead to less stable ink compositions.

Surprisingly, the parent compound, benzimidazole, is an effective inhibitor even when compared to the benzimidazole

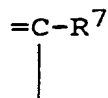
5 derivative which has been made and tested.

CLAIMS:

1. A printing ink composition comprising a resin, a solvent and an inhibitor; the inhibitor being benzimidazole or a compound having the general formula



wherein the A ring is benzenoid, naphthenoid or saturated cycloaliphatic, the A ring being unsubstituted or substituted with R^1 which is an alkyl group of 1 to 4 carbon atoms; and X is a nitrogen atom or the



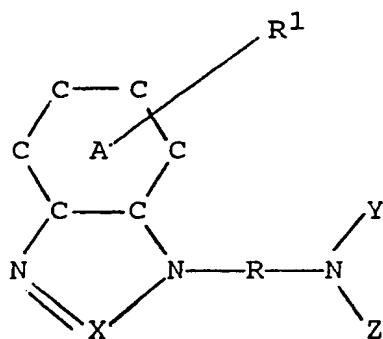
group, wherein R^7 is a hydrogen atom or an alkyl group of 1 to 4 carbon atoms; R is an alkylene group of 1 to 5 carbon atoms; Y is a hydrogen atom, a hydroxide group or an organic moiety and Z is an organic moiety or Y and Z when taken together with the nitrogen to which they are attached form an organic ring structure; the inhibitor when other than benzimidazole having a 24 hour room temperature isopropyl alcohol solubility of less than 5% by weight.

2. The ink composition of claim 1, wherein the inhibitor has a 24 hour room temperature aqueous solubility of less than 0.1% by weight.

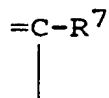
3. The ink composition of claim 1 or claim 2, wherein the inhibitor is N,N'-bis(tolyltriazol-1-yl methyl) piperazine, N,N'-bis(benzotriazol-1-yl methyl) piperazine, N,N'-bis(cyclohexyltriazol-1-yl methyl) piperazine, and N,N'-bis(methylcyclohexyltriazol-1-yl methyl) piperazine; 1,3-bis(benzotriazol-1'-yl methyl) urea; 1,3-bis(5'-tolyltriazol-1'-yl methyl) urea; 1,3-bis(benzotriazol-1'-yl methyl) N,N'-dimethyl urea; 1,5-bis(benzotriazol-1'-yl methyl) biuret; 2,4,6-tris(benzotriazol-1'-yl methyl)-s-triazine; 2,4-bis(benzotriazol-1'-yl methyl) benzoguanamine; 1-(1'-benzenesulfonamido) methyl benzotriazole; 1-(1'-methanesulfonamido)methyl benzotriazole; 4-(benzotriazol-1'-yl-methyl) hydantoin; 1-(1'-(2'-oxopyrrolidin-1'-yl) ethyl) benzotriazole; N,N-bis(benzotriazol-1-yl methyl) hydroxylamine; 1,3-bis(benzotriazol-1'-yl methyl) thiourea; N,N-bis(benzotriazol-1-yl methyl) glycine; N-(benzotriazol-1-yl methyl)-4'-carboxybenzene sulfonamide; N,N'-bis(benzotriazol-1-yl methyl) naphthalene-1,5-disulfonamide; N,N'-bis(benzotriazol-1-yl methyl) benzene-1,3-disulfonamide; 1-bis(benzotriazol-1'-yl methyl)-2-benzoyl hydrazide; bis(benzotriazol-1-yl methyl) amine;

1,3-bis(5'-butyl benzotriazol-1'-yl methyl) urea; or
 1,3-bis(benzimidazol-1'-yl methyl) piperazine.

4. The ink composition of claim 1 or claim 2,
 wherein the inhibitor is a compound having the general
 formula



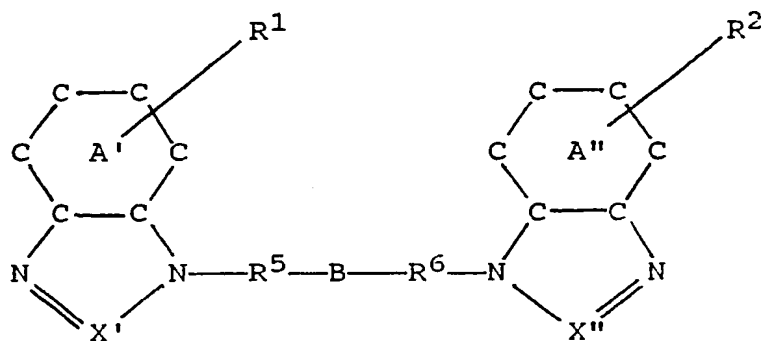
wherein the A ring is benzenoid, naphthenoid or saturated
 cycloaliphatic, the A ring being unsubstituted or
 substituted with R^1 which is an alkyl group of 1 to 4
 carbon atoms; X is a nitrogen atom or the



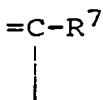
group, wherein R^7 is a hydrogen atom or an alkyl group of
 1 to 4 carbon atoms; R is an alkylene group of 1 to 5
 carbon atoms; Y is a hydrogen atom, an alkyl group of 1
 to 4 carbon atoms, an hydroxyl moiety or a carboxy
 containing moiety; and Z is methyl benzotriazole, methyl
 tolyltriazole, a carbonyl or thiocarbonyl linked methyl
 benzotriazole or methyl tolyltriazole containing moiety,
 or a sulfonyl linked moiety; or Y and Z taken together
 form a saturated ring compound containing a carbonyl

group, or having a methyl benzotriazole or methyl tolyltriazole containing moiety attached thereto.

5. The ink composition of claim 1 or claim 2, wherein the inhibitor is a compound having the formula



wherein each of the A' and A'' rings, which may be the same or different, is a benzenoid, naphthenoid or saturated cycloaliphatic moiety and are each unsubstituted or substituted with R¹ or R², which may be the same or different and each represents an alkyl group of 1 to 4 carbon atoms; R⁵ and R⁶, which may be the same or different, each represents an alkylene group of 1 to 5 carbon atoms, X' and X'', which may be the same or different, each represents a nitrogen atom or the



group wherein R⁷ is a hydrogen atom or an alkyl group of 1 to 4 carbon atoms; B is NH, NOH or an organic moiety; and both the R⁵-B and B-R⁶ bonds are carbon/nitrogen bonds.

6. The ink composition of claim 5, wherein X' and X'' are nitrogen atoms.

7. The ink composition of claim 5 or claim 6, wherein B is selected from the group consisting of organic substituted nitrogen, organic substituted nitrogen/carbonyl/nitrogen, organic substituted nitrogen/carbonyl/nitrogen/carbonyl/nitrogen, and organic substituted or unsubstituted saturated heterocyclic ring.

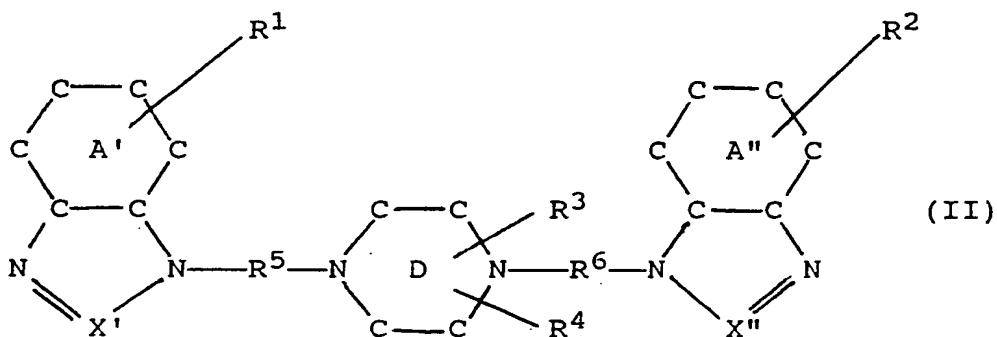
8. The ink composition of claim 1 or claim 2, wherein the inhibitor is a compound having at least two moieties selected from the group consisting of 1-methyl benzotriazole moiety, 1-ethyl benzotriazole moiety, 1-methyl benzimidazole moiety, 1-ethyl benzimidazole moiety, 1-methyl tolyltriazole moiety and 1-ethyl tolyltriazole moiety, the selected moieties being attached to one or more nitrogen atoms.

9. The ink composition of claim 1 or claim 2, wherein the inhibitor is a compound having at least one moiety selected from the group consisting of 1-methyl benzotriazole moiety, 1-ethyl benzotriazole moiety, 1-methyl tolyltriazole moiety and 1-ethyl tolyltriazole moiety, the selected moiety being attached to a nitrogen atom, and the inhibitor being a compound having at least one carboxy moiety.

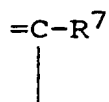
10. The ink composition of claim 1 or claim 2, wherein the inhibitor is a compound having at least one moiety selected from the group consisting of 1-methyl benzotriazole moiety, 1-ethyl benzotriazole moiety, 1-methyl tolyltriazole moiety and 1-ethyl tolyltriazole moiety, the selected moiety being attached to a nitrogen atom; and the inhibitor being a compound having at least one sulfonyl moiety attached to a nitrogen atom.

11. The ink composition of any one of claims 1 to 10, wherein the solvent is water.

12. A compound of the formula



wherein X' and X'', which may be the same or different, each represents a nitrogen atom or the



group wherein R⁷ is a hydrogen atom or an alkyl group of 1 to 4 carbon atoms, each of the A' and A'' rings which may be the same or different, is a benzenoid, saturated cycloaliphatic or naphthenoid moiety, each of R¹ and R²,

which may be the same or different, (a) when attached to a benzenoid moiety, represents a hydrogen atom or an alkyl group of 2 to 4 carbon atoms and (b) when attached to a saturated cycloaliphatic or naphthenoid moiety represents a hydrogen atom or an alkyl group of 1 to 4 carbon atoms, R^3 and R^4 , which may be the same or different, each represents a hydrogen atom or an alkyl group of 1 to 4 carbon atoms, R^5 and R^6 , which may be the same or different, each represents an alkylene group of 1 to 5 carbon atoms and the D ring is saturated; or the compound is selected from the group consisting of 1,3-bis(5'-tolyltriazol-1'-yl methyl) urea, 1,5-bis(benzotriazol-1'-yl methyl) biuret, 2,4,6-tris(benzotriazol-1'-yl methyl)-s-triazine, 2,4-bis(benzotriazol-1'-yl methyl) benzoguanamine, N,N-bis(benzotriazol-1-yl methyl) glycine, N-(benzotriazol-1-yl methyl)-4'-carboxybenzene sulfonamide, N,N'-bis(benzotriazol-1-yl methyl) naphthalene-1,5-disulfonamide, N,N'-bis(benzotriazol-1-yl methyl) benzene-1,3-disulfonamide, 1-bis(benzotriazol-1'-yl methyl)-2-benzoyl hydrazide, bis(benzotriazol-1-yl methyl) amine and 1,3-bis(5'-butyl benzotriazol-1'-yl methyl) urea.

Patents Act 1977
Examiner's report to the Comptroller under Section 17
(The Search report)

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Relevant Technical Fields

(i) UK Cl (Ed.N) C3V (VAD, VAE); C3K (KXX)

(ii) Int Cl (Ed.6) C09D

Search Examiner
K MACDONALD

Date of completion of Search
19 JULY 1995

Databases (see below)

(i) UK Patent Office collections of GB, EP, WO and US patent specifications.

(ii) ONLINE: WPI

Documents considered relevant
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Claims :-
1-11

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